

AMENDMENTS TO THE CLAIMS

This listing of claims shall replace all prior version and listings of claims in the application. Please amend the claims as follows:

Claim 1 (Currently amended): A method of assessing *de novo* fatty acid synthesis in a tissue of an organism, comprising quantifying a marker of *de novo* fatty acid synthesis in a biological fluid sample from the organism, and correlating the quantity of the marker to the level of *de novo* fatty acid synthesis in the tissue, wherein the biological fluid sample is [[a]] blood or a blood product, and wherein the marker of *de novo* fatty acid synthesis comprises:

- (a) palmitoleic acid or both the ratio of palmitoleic acid and to palmitic acid quantified from the free fatty acid fraction of the blood or blood product, wherein the method is a method to assess *de novo* fatty acid synthesis in adipose tissue; or
- (b) palmitoleic acid or both the ratio of palmitoleic acid and to palmitic acid quantified from the phosphatidylcholine or cholesterol ester fraction of the blood or blood product, wherein the method is a method to assess *de novo* fatty acid synthesis in liver tissue.

Claims 2-7 (Canceled)

Claim 8 (Currently amended): The method of claim 1, wherein the biological fluid sample is a plasma sample.

Claim 9 (Canceled)

Claim 10 (Withdrawn-currently amended): The method of claim 1 wherein the marker of *de novo* fatty acid synthesis is quantified from the free fatty acid fraction of the blood or blood product and the method is a method to assess *de novo* fatty acid synthesis in adipose tissue.

Claim 11 (Currently amended): The method of claim 1 wherein the marker of *de novo* fatty acid synthesis is quantified from the phosphatidylcholine or cholesterol ester fraction of the blood or blood product, and the method is a method to assess *de novo* fatty acid synthesis in liver tissue.

Claim 12 (Currently amended): The method of claim 1, wherein the marker comprises the ratio of comprising quantifying both palmitoleic acid to and palmitic acid in a biological fluid sample from the organism.

Claim 13 (Canceled)

Claim 14 (Currently amended): The method of claim [[13]] 12, wherein the correlating step comprises further comprising comparing the ratio of palmitoleic acid to palmitic acid in a indicator from the biological fluid sample with a ratio indicator from a baseline or control fluid sample.

Claims 15-20 (Canceled)

Claim 21 (Currently amended): The method of claim 1, wherein the method is

(1) a method to determine if a pharmaceutical, nutritional, genetic, toxicological or environmental treatment, regimen or dosage influences *de novo* fatty acid synthesis; or

(2) a method to assess a therapeutic or pharmaceutical agent for its potential effectiveness, efficacy or side effects relating to *de novo* fatty acid synthesis[[; or]],

(3) a method to screen individuals for compatibility or incompatibility with a pharmaceutical, nutritional, toxicological or environmental treatment.

Claim 22 (Currently amended): The method of claim 1, comprising quantifying wherein the marker of *de novo* fatty acid synthesis is palmitoleic acid in a biological sample from the organism.

Claims 23-25 (Canceled)

Claim 26 (Currently amended): The method of claim 1, wherein the method is a method of assessing a change in the *de novo* fatty acid synthesis in the organism, and wherein the method

comprises taking at least two biological fluid samples from the organism, wherein the two fluid samples are taken before and after an event.

Claim 27 (Original): The method of claim 26, wherein the event comprises passage of time, treatment with a therapeutic agent, treatment with a pharmaceutical agent, treatment with a nutritional regimen, treatment with a genetic modification, exposure to a toxic or potentially toxic compound, exposure to an environmental condition, treatment with a laboratory procedure, exercise, or the appearance of a phenotypic state.

Claims 28-31 (Canceled)

Claim 32 (Currently amended): The method of claim [[28]] 27, further comprising comparing the assessment of *de novo* fatty acid synthesis from the organism to an assessment of *de novo* fatty acid synthesis from another organism or compiled for a population of organisms.

Claim 33 (Currently amended): The method of claim [[28]] 27, wherein the quantity of the marker of *de novo* fatty acid synthesis is reported as an absolute or relative concentration.

Claim 34 (Original): The method of claim 33, wherein correlating the quantity of the marker of *de novo* fatty acid synthesis comprises using the absolute or relative concentration of the marker of *de novo* fatty acid synthesis in a mathematical or statistical equation for determining the amount of *de novo* fatty acid synthesis.

Claims 35-60 (Canceled)

Claim 61 (Currently amended): The method of claim 1, wherein the method is a method of assessing an activity of at least one enzyme involved in *de novo* fatty acid synthesis, further comprising correlating the quantity of the marker with the activity of the at least one enzyme.

Claim 62 (Original): The method of claim 1, further comprising generating a printed report.

Claim 63 (Currently amended): The method of claim 11, wherein the marker of *de novo* fatty acid synthesis is quantified from the cholesterol ester fraction of the blood or blood product.

Claim 64 (New): The method of claim 61, wherein the enzyme involved in *de novo* fatty acid synthesis is fatty acid synthase.

Claim 65 (New): The method of claim 61, wherein the enzyme involved in *de novo* fatty acid synthesis is stearoyl Coenzyme A-desaturase.

Claim 66 (New): The method of claim 1, wherein the marker of *de novo* fatty acid synthesis is quantified from the cholesterol ester fraction of the blood or blood product.